

IN THE CLAIMS

Please amend the claims as follows:

1. (Currently Amended) A multilayer dosage form comprising
  - a) a neutral core,
  - b) an inner methacrylate copolymer coating comprising at least 90% by weight of (meth)acrylate monomers having neutral radicals, wherein the methacrylate copolymer has a minimum film-forming temperature as specified in DIN 53 787 not exceeding 30°C, and a pharmaceutically active substance bound to the methacrylate polymer
  - c) an outer coating which comprises a (meth)acrylate copolymer which is composed of 10 to 30% by weight methyl methacrylate, 50 to 70% by weight methyl acrylate and 5 to 15% by weight methacrylic acid of a copolymer which is comprised of 40 to 95% by weight free radical polymerized C<sub>4</sub>-to C<sub>4</sub>-alkyl esters of acrylic or of methacrylic acid and 5 to 60% by weight (meth)acrylate monomers having an anionic group in the alkyl radical,  
wherein the values for the percentage release of active substance in a hypotonic and an isotonic release medium based on phosphate buffer pH 6.8 do not differ from one another at any time in the period from 1 to 5 hours by more than 10%.
2. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the methacrylate copolymer of the inner coating is polymerized from 25-35% by weight methyl methacrylate, 75 to 65% by weight ethyl acrylate and, optionally, up to 10% by weight other vinylically polymerizable monomers, wherein the proportionate amounts add up to 100% by weight.
3. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the active substance/polymer ratio of the inner layer is from 20:1 to 1:20.

4. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the outer coating comprises a (meth)acrylate copolymer which is composed of 40 to 60% by weight methacrylic acid and 60 to 40% by weight methyl methacrylate or 40 to 60% by weight methacrylic acid and 60 to 40% by weight ethyl acrylate.

5. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the outer coating comprises a (meth)acrylate copolymer which is composed of 20 to 40% by weight methacrylic acid and 80 to 60% by weight methyl methacrylate.

6. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the outer coating comprises a (meth)acrylate copolymer which is composed of 20 to 34% by weight methacrylic acid and/or acrylic acid, 20 to 69% by weight methyl acrylate, 0 to 40% by weight ethyl acrylate and, where appropriate, 0 to 10% by weight further vinylically copolymerizable monomers, wherein the glass transition temperature of the copolymer as specified in ISO 11357-2, subsection 3.3.3, does not exceed 60°C.

7. (Cancelled).

8. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein said multilayer dosage form comprises an active substance from the active substance classes of aminosalicylates, of sulfonamides or of glucocorticoids.

9. (Previously Presented) The multilayer dosage form as claimed in claim 8, wherein said multilayer dosage form comprises the active substance 5-aminosalicylic acid, olsalazine, sulfalazine, prednisone, prednisolone or budesonide.

10. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein said multilayer dosage form comprises an active substance from the active substance classes of enzymes, peptide hormones, immunomodulatory proteins, antigens, antibodies or of oligonucleotides.

11. (Previously Presented) The multilayer dosage form as claimed in claim 10, wherein said multilayer dosage form comprises the active substance pancreatin, insulin, human growth hormone (hGH), carboplatin, intron A, calcitonin, cromalyn, interferons, calcitonin, granulocyte colony stimulating factor (G-CSF), interleukin, parathyroid hormones, glucagon, pro-somatostatin, somatostatin, detirelix, cetorelix, vasopressin, 1-deaminocysteine-8-D-arginine-vasopressin, leuprolide acetate or an antigen which has been isolated from one or more grasses or one or more other plants.

12. (Cancelled).

13. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the inner coating comprises not more than 1% by weight of a release agent.

14. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the inner coating does not comprise release agents.

15. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the inner coating does not comprise plasticizer.

16. (Previously Presented) The multilayer dosage form as claimed in claim 1, wherein the methacrylate copolymer comprise 65% by ethyl acrylate and 35% by weight methyl methacrylate.